## Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application.

## Listing of Claims

## 1. (Currently amended) A compound of the formula

wherein D, E, F, G, L, T, W, X, Y and Z are each, independently, N or CH;

U is N and U is single bonded to both carbons adjacent to it in the nitrogen containing ring of which it is a member;

A is (CH<sub>2</sub>)<sub>m</sub> wherein m is zero, one or two;

R<sup>1</sup> and R<sup>2</sup> are selected, independently, from hydrogen, (C<sub>1</sub>-C<sub>6</sub>) alkyl optionally substituted with from one to seven fluorine atoms, cyano, -OR<sup>9</sup>, and -CONHR<sup>10</sup>;

or R<sup>1</sup> and R<sup>2</sup>, together with carbon atoms of the cyclopropyl ring to which they are attached, form a five or six membered saturated or unsaturated monocyclic ring containing from zero to four heteroatoms, wherein said heteroatoms are selected, independently, from oxygen, sulfur and nitrogen, with the provise that there can not be two adjacent ring oxygen atoms; and wherein said ring can be optionally substituted with from one to three substituents independently selected from (C<sub>1</sub> C<sub>4</sub>) alkyl optionally substituted with from one to three fluorine atoms, eyano, nitro, halo, hydroxy, amino, (C<sub>1</sub> C<sub>4</sub>) alkylamino, di[(C<sub>1</sub> C<sub>6</sub>)alkyl] amino, (C<sub>1</sub> C<sub>4</sub>) alkoxyl;

or one of R<sup>1</sup>-and R<sup>2</sup>-forms, together with R<sup>2</sup>, a five or six membered saturated or unsaturated monocyclic ring containing from zero to four heteratoms, wherein said heteratoms are selected, independently, from oxygen, sulfur and nitrogen, with the provise that there can not be two adjacent ring exygen atoms, and wherein said ring can be optionally substituted with from one to three substituents independently selected from (C<sub>1</sub>-C<sub>4</sub>) alkyl optionally substituted with

from one to three fluorine atoms, cyano, nitro, halo, hydroxy, amino, (C<sub>1</sub>-C<sub>4</sub>) alkylamino, di[(C<sub>1</sub>-C<sub>6</sub>)alkyl] amino, (C<sub>1</sub>-C<sub>4</sub>) amideamine and (C<sub>1</sub>-C<sub>4</sub>) alkanoyl;

R<sup>3</sup> and R<sup>4</sup> are selected, independently, from hydrogen, halo, (C<sub>1</sub>-C<sub>6</sub>) alkyl optionally substituted with from one to seven fluorine atoms, cyano, hydroxy, -CONHR<sup>11</sup>, -OR<sup>12</sup>, -NR<sup>13</sup>R<sup>14</sup> and -COR<sup>15</sup>:

or one of  $\mathbb{R}^3$  and  $\mathbb{R}^4$  forms, together with  $\mathbb{R}^7$ , a five-or-six membered aromatic or nonaromatic ring containing from one to-four hoteroatoms, wherein said heteroatoms are selected, independently, from oxygen, sulfur and nitrogen, with the provise that there can not be two adjacent ring oxygen atoms, and wherein said ring can be optionally substituted with from one-to-three substituents independently selected from  $(C_1, C_4)$  alkyl-optionally substituted with from one-to-three fluorine atoms,  $(C_1, C_4)$  alkoxy optionally substituted with from one-to-three fluorine atoms, eyano, nitro, halo, hydroxy, amino,  $(C_1, C_4)$  alkylamino, di[ $(C_1, C_5)$  alkyl-amino,  $(C_1, C_4)$  amidoamino and  $(C_1, C_4)$  alkanoyl;

R<sup>5</sup> and R<sup>6</sup> are selected, independently, from hydrogen, halo, (C<sub>1</sub>-C<sub>6</sub>) alkyl optionally substituted with from one to seven chlorine atoms, cyano, hydroxy, -CONHR<sup>16</sup>, -OR<sup>17</sup>, -NR<sup>18</sup>R<sup>19</sup>, and -COR<sup>20</sup>;

 $R^7$  is hydrogen, (C<sub>1</sub>-C<sub>6</sub>)alkyl optionally substituted with from one to seven fluorine atoms, or aryl selected from phenyl and naphthyl, wherein said aryl can be optionally substituted with from one to three substituents independently selected from (C<sub>1</sub>-C<sub>4</sub>) alkyl optionally substituted with from one to three fluorine atoms, (C<sub>1</sub>-C<sub>4</sub>) alkoxy optionally substituted with from one to three fluorine atoms, cyano, nitro, halo, hydroxy, amino, (C<sub>1</sub>-C<sub>4</sub>) alkylamino, di[(C<sub>1</sub>-C<sub>6</sub>)alkyl] amino, (C<sub>1</sub>-C<sub>4</sub>) amidoamino and (C<sub>1</sub>-C<sub>4</sub>) alkanoyl;

or  $\mathbb{R}^3$ -can form a ring with  $\mathbb{R}^4$ -or  $\mathbb{R}^2$ , as described in the above definition of  $\mathbb{R}^4$  and  $\mathbb{R}^2$ ; or  $\mathbb{R}^2$ -can form a ring with  $\mathbb{R}^3$ -or  $\mathbb{R}^4$ , as described in the above definition of  $\mathbb{R}^3$ -and  $\mathbb{R}^4$ ;  $\mathbb{R}^8$  is selected from hydrogen, cyano,  $(C_1-C_6)$  alkyl optionally substituted with from one to seven fluorine atoms,  $-O\mathbb{R}^9$ , and  $-CONH\mathbb{R}^{10}$ ;

R<sup>9</sup>, R<sup>10</sup>, R<sup>11</sup>, R<sup>12</sup>, R<sup>13</sup>, R<sup>14</sup>, R<sup>16</sup>, R<sup>17</sup>, R<sup>18</sup> and R<sup>19</sup> are selected, independently, from hydrogen, (C<sub>1</sub>-C<sub>6</sub>)alkyl optionally substituted with from one to seven fluorine atoms, aryl and heteroaryl, wherein said aryl is selected from phenyl and naphthyl and said heteroaryl is selected from four to six membered monocyclic aromatic rings containing from one to four heteroatoms (nonlimiting examples of such rings are furyl, thienyl, pyrrolyl, oxazelyl, thiazolyl, imidazelyl, oxadiazelyl, thiadiazelyl, pyridyl, triazelyl, triazelyl, triazelyl, pyridazyl,

pyrimidinyl and pyrazolyl) and eight to twelve membered bicyclic aromatic rings containing from one to five heteroatoms, wherein said heteroatoms are selected, independently, from oxygen, sulfur and nitrogen, with the proviso that there can not be two adjacent ring oxygen atoms, and wherein said aryl and heteroaryl rings can optionally be substituted one or more substitutents, preferably with from zero to two substituents, independently selected from  $(C_1-C_4)$  alkyl optionally substituted with from one to three fluorine atoms,  $(C_1-C_4)$  alkoxy optionally substituted with from one to three fluorine atoms, cyano, nitro, halo, hydroxy, amino,  $(C_1-C_4)$  alkylamino, di[ $(C_1-C_6)$ alkyl] amino,  $(C_1-C_4)$  amidoamino and  $(C_1-C_4)$  alkanoyl;

R<sup>15</sup> and R<sup>20</sup> are selected, independently, from NHR<sup>21</sup> and the group of radicals listed in the definition of R<sup>9</sup> through R<sup>19</sup>; and

R<sup>21</sup> is selected from the group of radicals listed in the definition of R<sup>9</sup> through R<sup>19</sup>;

or a pharmaceutically acceptable salt thereof, with the proviso that ring D, E, F, G, L and ring T, W, X, Y, and Z are each independently pyrimidyl, or phenyl rings.

- 2. (Cancelled).
- 3. (Original) A compound according to claim 1, wherein both R<sup>1</sup> and R<sup>2</sup> are hydrogen.
- 4. (Original) A compound according to claim 1, wherein  $R^1$  and  $R^2$  are selected, independently, from hydrogen and  $(C_1 C_6)$  alkyl.
  - 5-11. (Cancelled)
- 12. (Original) A compound according to claim 1 which is in the Z (cis) configuration with respect to the cyclopropyl ring.
- 13. (Currently amended) A compound according to claim 1 wherein m is zero or two, in the case where m is two forming an azabicyclic ring system bridged either diagonally or directly across the ring system the compound is:

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2,6-dimethyl-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-4-chloro-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2,4,6-trimethyl-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-3,4-dimethyl-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-3,5-dimethoxy-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-3-fluoro-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2-methyl-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2,3-dimethyl-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl}-cyclopropylmethyl}-3,4-dimethoxy-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl}-cyclopropylmethyl}-3,4,5-trimethoxy-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-4-fluoro-benzamide:

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-vl)-piperazin-1-ylmethyl}-cyclopropylmethyl}-3-chloro-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-vl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-4-trifluoromethyl-benzamide:

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-3-cyano-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2,6-dimethyl-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2,4-difluoro-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2,3-difluoro-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2-trifluoromethyl-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl}-cyclopropylmethyl}-2,5-dichloro-benzamide;

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[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2,3-dichloro-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2,4-dimethyl-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-3-chloro-2-fluoro-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-2-chloro-4-fluoro-benzamide;

[Z](+/-)N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl}-cyclopropylmethyl}-2,4-dichloro-benzamide; and

N-{2-[4-(2-tert-Butyl-6-trifluoromethyl-pyrimidin-4-yl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-4-diethylamino-benzamide.

- 14. (Original) A compound according to claim 1 wherein R<sup>1</sup> and R<sup>2</sup> are selected, independently, from hydrogen, methyl, cyano, trifluoromethyl and trifluoromethoxy.
- 15. (Currently amended) A pharmaceutical composition for treating a disorder or condition selected from psychotic disorders (e.g., psychosis, schizophrenia, schizo affective disorders, psychotic depression, mania, paranoid and delusional disorders), anxiety related disorders (e.g., generalized anxiety disorder, post traumatic stress disorder, panie disorder, obsessive compulsive disorder and phobias, including social phobia), mood disorders (e.g., cyclothymia, dysthymia, major depressive disorder, premenstrual syndrome, premenstrual dysphorie-disorder, bipolar disorder, seasonal affective disorder), and Parkinson's disease, hypertension, hypotension, urinary incontinence, chemical dependencies and addictions (e.g., dependencies on alcohol, cocaine, heroin, nicotino, benzodiazepines, phenobarbitel), sexual dysfunction (e.g., premature ejaculation, male erectile dysfunction) and movement disorders (e.g., drug induced and neurodegeneration based dyskinesias) in a mammal, comprising an amount of a compound according to claim 1, or a pharmaceutically acceptable salt thereof, that is effective in treating such disorder or condition, and a pharmaceutically acceptable carrier.
- 16. (Currently amended) A method of treating a disorder or condition selected from psychotic <u>disorders(e.g., psychosis, schizophrenia, schizo affective disorders, psychotic depression, mania, paranoid and delusional disorders), anxiety related disorders (e.g.,</u>

generalized anxiety disorder, post traumatic stress disorder, panic disorder, obsessivecompulsive disorder and phobias, including social phobia), mood disorders (e.g.,
eyelothymia, dysthymia, major depressive disorder, premenstrual syndrome, premenstrual
dysphoric disorder, bipolar disorder, seasonal affective disorder), and Parkinson's disease,
hypertension, hypotension, urinary incontinence, chemical dependencies and addictions (e.g.,
dependencies on alcohol, cocaine, heroin, nicotine, benzodiazopines, phenobarbitol), sexual
dysfunction (e.g., premature ejaculation, male erectile dysfunction) and movement disorders
(e.g., drug induced and neurodegeneration based dyskinesias)- in a mammal, comprising
administering to said mammal an amount of a compound according to claim 1, or a
pharmaceutically acceptable salt thereof, that is effective in treating such disorder or
condition.

17.-22. (Cancelled)

23. (Original) A method according to claim 16 wherein the disorder or condition being treated is a psychotic disorder or condition.

24.-25. (Cancelled)

26. (Original) A method according to claim 16 wherein the disorder or condition being treated is Parkinson's disease.

27.-32. (Cancelled)

33. (New) A compound according to claim 1 wherein the compound is: [Z](+/-)2,6-Dimethyl-N-{2-[4-(3-trifluoromethyl-phenyl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-benzamide; and [E](+/-)2,6-Dimethyl-N-{2-[4-(3-trifluoromethyl-phenyl)-piperazin-1-ylmethyl]-cyclopropylmethyl}-benzamide.